

CLAIMS

1. The use of an anti-angiogenic agent in combination with an inhibitor of the Src family of non-receptor tyrosine kinases in the manufacture of a medicament for use in the substantially normotensive treatment in a warm-blooded mammal such as a human being of a disease state associated with angiogenesis, the Src kinase inhibitor being administered in an amount effective to counteract substantially the hypertension induced by the anti-angiogenic agent.
2. The use of an anti-angiogenic agent in combination with a Src kinase inhibitor according to claim 1 in the manufacture of a medicament for use in the substantially normotensive production of an anti-cancer effect in a warm-blooded mammal such as a human being, the Src kinase inhibitor being administered in an amount effective to counteract substantially the hypertension induced by the anti-angiogenic agent.
3. The use according to claim 1 of an anti-angiogenic agent in combination with a Src kinase inhibitor wherein the anti-angiogenic agent is selected from :-
 - 4-(4-fluoro-2-methylindol-5-yloxy)-6-methoxy-7-(3-(pyrrolidin-1-yl)propoxy)quinazoline,
 - 4-(4-fluoro-2-methylindol-5-yloxy)-6-methoxy-7-(3-piperidinopropoxy)quinazoline,
 - 4-(4-fluoro-2-methylindol-5-yloxy)-6-methoxy-7-((1-methylpiperidin-4-yl)methoxy)quinazoline,
 - 4-(4-fluoro-2-methylindol-5-yloxy)-6-methoxy-7-(3-(4-methylpiperazin-1-yl)propoxy)quinazoline,
 - 4-(4-fluoro-2-methylindol-5-yloxy)-6-methoxy-7-(2-(1-methylpiperidin-4-yl)ethoxy)quinazoline,
 - 4-(4-chloro-2-fluoroanilino)-6-methoxy-7-(1-methylpiperidin-4-ylmethoxy)quinazoline,
 - 4-(4-bromo-2-fluoroanilino)-6-methoxy-7-(1-methylpiperidin-4-ylmethoxy)quinazoline,
 - 4-(4-chloro-2-fluoroanilino)-6-methoxy-7-(piperidin-4-ylmethoxy)quinazoline,
 - 4-(4-bromo-2-fluoroanilino)-6-methoxy-7-(piperidin-4-ylmethoxy)quinazoline,
 - 4-[(4-fluoro-2-methyl-1*H*-indol-5-yl)oxy]-6-methoxy-7-{3-[4-(2,2,2-trifluoroethyl)piperazin-1-yl]propoxy}quinazoline,
 - 7-{2-[4-(2-fluoroethyl)piperazin-1-yl]ethoxy}-4-[(4-fluoro-2-methyl-1*H*-indol-5-yl)oxy]-6-methoxyquinazoline,

- 4-[(4-fluoro-2-methylindol-5-yl)oxy]-6-methoxy-7-{3-[4-(2-propynyl)piperazin-1-yl]propoxy}quinazoline,
 7-{3-[4-(2-fluoroethyl)piperazin-1-yl]propoxy}-4-[(4-fluoro-2-methyl-1*H*-indol-5-yl)oxy]-6-methoxyquinazoline,
 5 7-(3-(4-acetylpiperazin-1-yl)propoxy)-4-(4-fluoro-2-methylindol-5-yloxy)-6-methoxyquinazoline, and
 7-[2-(4-acetylpiperazin-1-yl)ethoxy]-4-[(4-fluoro-2-methyl-1*H*-indol-5-yl)oxy]-6-methoxyquinazoline;
 or a pharmaceutically-acceptable acid-addition salt thereof.

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4. The use according to claim 1 of an anti-angiogenic agent in combination with a Src kinase inhibitor wherein the Src kinase inhibitor selected from :-
 4-(2,4-dichloro-5-methoxyanilino)-7-(2-piperidinoethoxy)-5-tetrahydropyran-4-yloxyquinazoline,
 15 4-(2,4-dichloro-5-methoxyanilino)-7-(2-morpholinoethoxy)-5-tetrahydropyran-4-yloxyquinazoline,
 4-(6-chloro-2,3-methylenedioxyanilino)-7-(2-pyrrolidin-1-ylethoxy)-5-tetrahydropyran-4-yloxyquinazoline,
 4-(6-chloro-2,3-methylenedioxyanilino)-7-(3-pyrrolidin-1-ylpropoxy)-5-tetrahydropyran-4-yloxyquinazoline,
 20 4-(6-chloro-2,3-methylenedioxyanilino)-7-[2-(4-methylpiperazin-1-yl)ethoxy]-5-tetrahydropyran-4-yloxyquinazoline,
 4-(6-chloro-2,3-methylenedioxyanilino)-7-(2-piperidinoethoxy)-5-tetrahydropyran-4-yloxyquinazoline,
 25 7-[2-(4-acetylpiperazin-1-yl)ethoxy]-4-(6-chloro-2,3-methylenedioxyanilino)-5-isopropoxyquinazoline,
 6-methoxy-4-(2,3-methylenedioxyanilino)-7-(3-morpholinopropoxy)quinazoline,
 6-methoxy-4-(2,3-methylenedioxyanilino)-7-(3-pyrrolidin-1-ylpropoxy)quinazoline,
 6-methoxy-4-(2,3-methylenedioxyanilino)-7-[3-(4-methylpiperazin-1-yl)propoxy]quinazoline,
 30 6-methoxy-4-(2,3-methylenedioxyanilino)-7-(3-piperidinopropoxy)quinazoline,
 4-(6-chloro-2,3-methylenedioxyanilino)-7-[3-(4-isobutyrylpiperazin-1-yl)propoxy]-6-methoxyquinazoline,
 4-(2-chloro-5-methoxyanilino)-6-methoxy-7-(*N*-methylpiperidin-4-ylmethoxy)quinazoline,

- 4-(2-chloro-5-methoxyanilino)-6-methoxy-7-piperidin-4-ylmethoxyquinazoline,
 4-(2,4-dichloro-5-methoxyanilino)-6-methoxy-7-(N-methylpiperidin-
 4-ylmethoxy)quinazoline,
 4-(2,4-dichloro-5-methoxyanilino)-6-methoxy-7-piperidin-4-ylmethoxyquinazoline,
 5 7-[2-(4-acetylpiperazin-1-yl)ethoxy]-4-(5-chloro-2,3-methylenedioxyppyrid-4-ylamino)-
 5-tetrahydropyran-4-yloxyquinazoline,
 4-(5-chloro-2,3-methylenedioxyppyrid-4-ylamino)-7-{2-[(3RS,4SR)-
 3,4-methylenedioxyppyrolidin-1-yl)ethoxy]-5-tetrahydropyran-4-yloxyquinazoline,
 7-[2-(4-acetylpiperazin-1-yl)ethoxy]-4-(5-chloro-2,3-methylenedioxyppyrid-4-ylamino)-
 10 5-isopropoxyquinazoline and
 4-(5-chloro-2,3-methylenedioxyppyrid-4-ylamino)-
 7-{2-[(3RS,4SR)-3,4-methylenedioxyppyrolidin-1-yl)ethoxy]-5-isopropoxyquinazoline;
 or a pharmaceutically-acceptable acid-addition salt thereof.
- 15 5. A synergistic combination product comprising an anti-angiogenic agent and a
 Src kinase inhibitor for use simultaneously, sequentially or separately in the production of an
 anti-cancer effect in a warm-blooded animal such as a human being.
6. A synergistic combination product according to claim 5 wherein the anti-angiogenic
 20 agent is selected from the compounds listed in claim 3 and the Src kinase inhibitor is selected
 from the compounds listed in claim 4.
7. A blood pressure effect sparing combination product comprising an anti-angiogenic
 agent and a Src kinase inhibitor for use simultaneously, sequentially or separately in the
 25 production of an anti-cancer effect in a warm-blooded animal such as a human being.
8. A blood pressure effect sparing combination according to claim 7 wherein the
 anti-angiogenic agent is selected from the compounds listed in claim 3 and the Src kinase
 inhibitor is selected from the compounds listed in claim 4.
- 30 9. A combination product comprising an anti-angiogenic agent and a Src kinase inhibitor
 for use simultaneously, sequentially or separately in the production of a substantially
 normotensive anti-cancer effect in a warm-blooded mammal such as a human being.

10. A combination product according to claim 9 wherein the an anti-angiogenic agent is selected from :-

- 4-(4-fluoro-2-methylindol-5-yloxy)-6-methoxy-7-(3-(pyrrolidin-1-yl)propoxy)quinazoline,
 5 4-(4-fluoro-2-methylindol-5-yloxy)-6-methoxy-7-(3-piperidinopropoxy)quinazoline,
 4-(4-fluoro-2-methylindol-5-yloxy)-6-methoxy-7-(3-(4-methylpiperazin-1-yl)propoxy)quinazoline,
 4-(4-fluoro-2-methylindol-5-yloxy)-6-methoxy-7-(2-(1-methylpiperidin-4-yl)ethoxy)quinazoline,
 10 4-(4-chloro-2-fluoroanilino)-6-methoxy-7-(1-methylpiperidin-4-ylmethoxy)quinazoline,
 4-(4-bromo-2-fluoroanilino)-6-methoxy-7-(1-methylpiperidin-4-ylmethoxy)quinazoline,
 4-(4-bromo-2-fluoroanilino)-6-methoxy-7-(piperidin-4-ylmethoxy)quinazoline,
 4-[(4-fluoro-2-methylindol-5-yl)oxy]-6-methoxy-7-{3-[4-(2-propynyl)piperazin-1-yl]propoxy}quinazoline,
 15 7-(3-(4-acetylpiperazin-1-yl)propoxy)-4-(4-fluoro-2-methylindol-5-yloxy)-6-methoxyquinazoline and
 7-[2-(4-acetylpiperazin-1-yl)ethoxy]-4-[(4-fluoro-2-methyl-1H-indol-5-yl)oxy]-6-methoxyquinazoline,
 or a pharmaceutically-acceptable acid-addition salt thereof;
 20 and the Src kinase inhibitor is selected from :-
 4-(6-chloro-2,3-methylenedioxyanilino)-7-(2-pyrrolidin-1-ylethoxy)-5-tetrahydropyran-4-yloxyquinazoline,
 4-(6-chloro-2,3-methylenedioxyanilino)-7-(3-pyrrolidin-1-ylpropoxy)-5-tetrahydropyran-4-yloxyquinazoline,
 25 4-(6-chloro-2,3-methylenedioxyanilino)-7-[2-(4-methylpiperazin-1-yl)ethoxy]-5-tetrahydropyran-4-yloxyquinazoline,
 4-(6-chloro-2,3-methylenedioxyanilino)-7-(2-piperidinoethoxy)-5-tetrahydropyran-4-yloxyquinazoline,
 7-[2-(4-acetylpiperazin-1-yl)ethoxy]-4-(6-chloro-2,3-methylenedioxyanilino)-
 30 5-isopropoxyquinazoline,
 6-methoxy-4-(2,3-methylenedioxyanilino)-7-(3-morpholinopropoxy)quinazoline,
 6-methoxy-4-(2,3-methylenedioxyanilino)-7-(3-piperidinopropoxy)quinazoline,

- 4-(6-chloro-2,3-methylenedioxyanilino)-7-[3-(4-isobutylpiperazin-1-yl)propoxy]-
6-methoxyquinazoline,
4-(2-chloro-5-methoxyanilino)-6-methoxy-7-(N-methylpiperidin-4-ylmethoxy)quinazoline,
7-[2-(4-acetylpiperazin-1-yl)ethoxy]-4-(5-chloro-2,3-methylenedioxy-4-ylamino)-
5 5-tetrahydropyran-4-yloxyquinazoline,
4-(5-chloro-2,3-methylenedioxy-4-ylamino)-7-{2-[(3RS,4SR)-
3,4-methylenedioxy-1-yl)ethoxy]-5-tetrahydropyran-4-yloxyquinazoline,
7-[2-(4-acetylpiperazin-1-yl)ethoxy]-4-(5-chloro-2,3-methylenedioxy-4-ylamino)-
5-isopropoxyquinazoline and
10 4-(5-chloro-2,3-methylenedioxy-4-ylamino)-
7-{2-[(3RS,4SR)-3,4-methylenedioxy-1-yl)ethoxy]-5-isopropoxyquinazoline,
or a pharmaceutically-acceptable acid-addition salt thereof.
11. A pharmaceutical composition comprising a combination product according to any of
15 claims 5 to 10 and a pharmaceutically-acceptable excipient or carrier.
12. A combination product according to any of claims 5 to 10 wherein the anti-angiogenic
agent is 4-(4-fluoro-2-methylindol-5-yloxy)-6-methoxy-7-(3-piperidinopropoxy)quinazoline,
or a pharmaceutically-acceptable acid-addition salt thereof, and the Src kinase inhibitor is
20 7-[2-(4-acetylpiperazin-1-yl)ethoxy]-4-(6-chloro-2,3-methylenedioxyanilino)-
5-isopropoxyquinazoline, or a pharmaceutically-acceptable acid-addition salt thereof.